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Amendment to the Claims

1-33. (canceled)

34. (currently amended) A composition comprising a CD20 binding molecule, wherein the CD20 binding molecule comprises: a) a light chain variable region and a heavy chain variable region, wherein:

the light chain variable region comprises:

i) a CDRL1 amino acid sequence selected from the group consisting of SEQ ID NO:1, SEQ ID NO:3, and SEQ ID NO:5;

ii) a CDRL2 amino acid sequence selected from the group consisting of SEQ ID NO:7, SEQ ID NO:11, and SEQ ID NO:13; and

-iii) a CDRL3 amino acid sequence selected from the group consisting of SEQ ID NO:17, SEQ ID NO:19, and SEQ ID NO:21;

-iv) an FRL1 amino acid sequence consisting of SEQ ID NO:71;

v) an FRL2 amino acid sequence consisting of SEQ ID NO:72;

- vi) an FRL3 amino acid sequence consisting of SEQ ID NO:73; and

- vii) an FRL4 amino acid sequence consisting of SEQ ID NO:74.

b) a said heavy chain variable region, wherein the heavy chain variable region comprises:

i) a CDRH1 amino acid sequence selected from the group consisting of SEQ ID NO:23 and SEQ ID NO:25;

—ii) a CDRH2 amino acid sequence selected from the group consisting of SEQ ID NO:27, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:35, SEQ ID NO:37, and SEQ ID NO:39; and

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—iii) a CDRH3 amino acid sequence selected from the group consisting of SEQ ID NO:43, SEQ ID NO:45, SEQ ID NO:47, SEQ ID NO:49, SEQ ID NO:51, SEQ ID NO:53, SEQ ID NO:55, and SEQ ID NO:57;

iv) an FRH1 amino acid sequence consisting of SEQ ID NO:79;

- -v) an FRH2 amino acid sequence consisting of SEQ ID NO:80;
- vi) an FRH3 amino acid sequence consisting of SEQ ID NO:81; and
- -vii) an FRH4 amino acid sequence consisting of SEQ ID NO:82.

35-39. (canceled)

40. (withdrawn-currently amended) A method of treating B cell lymphoma comprising administering to a subject a composition comprising a CD20 binding molecule, wherein the CD20 binding molecule comprises: a) a light chain variable region and a heavy chain variable region, wherein:

the light chain variable region comprises:

-i) a CDRL1 amino acid sequence selected from the group consisting of SEQ ID NO:1, SEQ ID NO:3, and SEQ ID NO:5;

ii) a CDRL2 amino acid sequence selected from the group consisting of SEQ ID NO:7, SEQ ID NO:11, and SEQ ID NO:13; and

-iii) a CDRL3 amino acid sequence selected from the group consisting of SEQ ID NO:17, SEQ ID NO:19, and SEQ ID NO:21;

-iv) an FRL1 amino acid sequence consisting of SEQ ID NO:71;

- v) an FRL2 amino acid sequence consisting of SEQ ID NO:72;
- -vi) an FRL3 amino acid sequence consisting of SEQ ID NO:73; and
- vii) an FRL4 amino acid sequence consisting of SEQ ID NO:74.

b) a said heavy chain variable region, wherein the heavy chain variable region comprises:

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-i) a CDRH1 amino acid sequence selected from the group consisting of SEQ ID NO:23 and SEQ ID NO:25;

—ii) a CDRH2 amino acid sequence selected from the group consisting of SEQ ID NO:27, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:35, SEQ ID NO:37, and SEQ ID NO:39; and

—iii) a CDRH3 amino acid sequence selected from the group consisting of SEQ ID NO:43, SEQ ID NO:45, SEQ ID NO:47, SEQ ID NO:49, SEQ ID NO:51, SEQ ID NO:53, SEQ ID NO:55, and SEQ ID NO:57;

- iv) an FRH1 amino acid sequence consisting of SEQ ID NO:79;
- -v) an FRH2 amino acid sequence consisting of SEQ ID NO:80;
- vi) an FRH3 amino acid sequence consisting of SEQ ID NO:81; and
 vii) an FRH4 amino acid sequence consisting of SEQ ID NO:82.
- 41. (withdrawn) The method of Claim 40, wherein the CD20 binding molecule comprises the AME 33 Fab.
- 42. (withdrawn) The method of Claim 40, wherein the CD20 binding molecule has a binding affinity (K_d) for human CD20 of 5.0 x 10^{-10} M or less, and a dissociation rate (koff) for human CD20 of 5.0 x 10^{-4} s⁻¹ or less.
- 43. (withdrawn) The method of Claim 42, wherein the CD20 binding molecule has a binding affinity (K_d) for human CD20 of 1.5 x 10^{-10} M or less.
- 44. (withdrawn) The method of Claim 42, wherein the CD20 binding molecule has a dissociation rate (k_{off}) for human CD20 of 2.5 x 10^{-4} s⁻¹ or less.
- 45. (withdrawn) The method of Claim 42, wherein the CD20 binding molecule has an association rate (k_{on}) for human CD20 of 5.0 x 10^{-5} M⁻¹ s⁻¹ or greater.
- 46. (withdrawn) The method of Claim 40, wherein the B cell lymphoma is Non-Hodgkin's lymphoma.

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- 47. (withdrawn) The method of Claim 46, wherein the Non-Hodgkin's lymphoma is Waldenstrom's macroglobulinemia.
- 48. (currently amended) A <u>The</u> composition of Claim 34, wherein the light chain variable region comprises an amino acid sequence of SEQ ID NO:59 and the heavy chain variable region comprises an amino acid sequence of SEQ ID NO:61.
- 49. (new) A composition comprising a CD20 binding molecule, wherein the CD20 binding molecule comprises a light chain variable region and a heavy chain variable region, wherein:

the light chain variable region comprises:

a CDRL1 amino acid sequence of SEQ ID NO:3;

a CDRL2 amino acid sequence of SEQ ID NO:11; and

a CDRL3 amino acid sequence SEQ ID NO:21, and

the heavy chain variable region comprises:

a CDRH1 amino acid sequence of SEQ ID NO:23;

a CDRH2 amino acid sequence of SEQ ID NO:31; and

a CDRH3 amino acid sequence of SEQ ID NO:45.

- 50. (new) The composition of Claim 49, wherein the light chain variable region comprises an amino acid sequence of SEQ ID NO:63 and the heavy chain variable region comprises an amino acid sequence of SEQ ID NO:65.
- 51. (new) A composition comprising a CD20 binding molecule, wherein the CD20 binding molecule comprises a light chain amino acid sequence of SEQ ID NO:67 and a heavy chain amino acid sequence of SEQ ID NO:69.